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This file contains CAS Registry Numbers for easy and accurate substance identification.

=> S POLYAMIDE

113812 POLYAMIDE

89335 POLYAMIDES

L1 145425 POLYAMIDE

(POLYAMIDE OR POLYAMIDES)

=> S CELLULOSE

310336 CELLULOSE

4035 CELLULOSES

L2 310796 CELLULOSE

(CELLULOSE OR CELLULOSES)

=> S BLEND OR MIXTURE

130337 BLEND

131052 BLENDS

179620 BLEND

(BLEND OR BLENDS)

64689 MIXTURE

122972 MIXTURES

184633 MIXTURE

(MIXTURE OR MIXTURES)

1379307 MIXT

509718 MIXTS

1701553 MIXT

(MIXT OR MIXTS)

1743586 MIXTURE

(MIXTURE OR MIXT)

L3 1877485 BLEND OR MIXTURE

=> S L1 ANDL2

MISSING OPERATOR L1 ANDL2

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=> S L1 AND L2

L4 8115 L1 AND L2

=> S L3 AND L4

L5 2362 L3 AND L4

=> S FABRIC(L)ARTICLES

90338 FABRIC

81955 FABRICS

124943 FABRIC (FABRIC OR FABRICS) 69892 ARTICLES 1453 FABRIC (L) ARTICLES L6 => S L5 AND L6 11 L5 AND L6 L7 => S DYES 194562 DYES L8 => S L7 AND L8 2 L7 AND L8 L9 => D L9 BIB, ABS ANSWER 1 OF 2 CAPLUS COPYRIGHT 2003 ACS on STN T.9 1962:483826 CAPLUS AN 57:83826 DΝ OREF 57:16798e-h Water-soluble dyes containing methylol groups TIWaechter, Rudolf; Weissauer, Hermann; Braun, Willy IN Badische Anilin-& Soda-Fabrik A.-G. PA5 pp.; Addn. to Brit. 842,802 (CA 55, 8872f) SO דת Patent Unavailable T.A APPLICATION NO. DATE PATENT NO. KIND DATE _____ ____ -----19620711 GB PΤ GB 900950 PRAI DE 19590609 Dyes of the formula (D) (CH2OH)n, in which D is a radical of an AB azo or anthraquinone dye contg. .gtoreq. one 2,4-diamino-1,3,5-triazine group and .gtoreq. one water-solubilizing group, and in which H atoms susceptible to reaction with HCHO have been replaced by n (1-8) methylol groups, are used to dye articles of natural, regenerated, or chem.-modified cellulose, of natural, regenerated, or synthetic polyamides, or of polyurethans. The dyes may be prepd. by treating a dye (parent of D) with alkali and aq. HCHO, or by treating intermediates for the production of dyes with HCHO and then converting the methylol compds. to dyes of the above formula. Thus, water 60, 30% aq. HCHO 80, Na2CO3 36, and a dye 38.2, obtained by conpling 2-H2NC6H4SO3H (I) 17.3 with 2-[(2-hydroxybenzylidene)amino]-4,6diamino-1,3,5-triazine 23 parts, are stirred for 24 hrs. at room temp. The mixt. is salted, filtered, the residue washed with dil. NaCl until neutral, and then dried at 45.degree. in vacuo. It dyes cotton reddish yellow shades of good wetfastness. Similarly, 2-[(4-aminophenylsulfonyl)amino]-4,6-diamino-1,3,5-triazine was diazotized, coupled with 1,3,5-HO(HO3S)2C10H5, and finally treated with aq. HCHO to give an orange-red dye. 1-Amino-4-[p-[(4,6-diamino-1,3,5triazin - 2 - yl)sulfamoyl] anilino)] - 2 - anthraquinonesulfonic acid, treated with aq. HCHO gave a compd. which dyed a mixed fabric of staple fiber and cotton a wetfast red shade. Diazotized 1,3,7-(HO3S)2(H2N)C10H5 coupled with 2,4-diamino-6-(4-hydroxyanilino)-1,3,5-triazine and treated with aq. HCHO gave a dye which dyed cotton a wetfast reddish yellow shade. => D L9 1-2 BIB, ABS ANSWER 1 OF 2 CAPLUS COPYRIGHT 2003 ACS on STN L9 1962:483826 CAPLUS AN

DN

57:83826 OREF 57:16798e-h

Water-soluble dyes containing methylol groups

Waechter, Rudolf; Weissauer, Hermann; Braun, Willy IN Badische Anilin-& Soda-Fabrik A.-G. PΑ 5 pp.; Addn. to Brit. 842,802 (CA 55, 8872f) SO DTPatent LΑ Unavailable PATENT NO. KIND DATE APPLICATION NO. DATE ~---_____ 19620711 GB 900950 PΙ 19590609 PRAI DE Dyes of the formula (D)(CH2OH)n, in which D is a radical of an azo or anthraquinone dye contg. .gtoreq. one 2,4-diamino-1,3,5-triazine group and .gtoreq. one water-solubilizing group, and in which H atoms susceptible to reaction with HCHO have been replaced by n (1-8) methylol groups, are used to dye articles of natural, regenerated, or chem.-modified cellulose, of natural, regenerated, or synthetic polyamides, or of polyurethans. The dyes may be prepd. by treating a dye (parent of D) with alkali and aq. HCHO, or by treating intermediates for the production of dyes with HCHO and then converting the methylol compds. to dyes of the above formula. Thus, water 60, 30% aq. HCHO 80, Na2CO3 36, and a dye 38.2, obtained by conpling 2-H2NC6H4SO3H (I) 17.3 with 2-[(2-hydroxybenzylidene)amino]-4,6diamino-1,3,5-triazine 23 parts, are stirred for 24 hrs. at room temp. The mixt. is salted, filtered, the residue washed with dil. NaCl until neutral, and then dried at 45.degree. in vacuo. It dyes cotton reddish yellow shades of good wetfastness. Similarly, 2-[(4-aminophenylsulfonyl)amino]-4,6-diamino-1,3,5-triazine was diazotized, coupled with 1,3,5-HO(HO3S)2C10H5, and finally treated with aq. HCHO to give an orange-red dye. 1-Amino-4-[p-[(4,6-diamino-1,3,5triazin - 2 - yl)sulfamoyl] anilino)] - 2 - anthraquinonesulfonic acid, treated with aq. HCHO gave a compd. which dyed a mixed fabric of staple fiber and cotton a wetfast red shade. Diazotized 1,3,7-(HO3S)2(H2N)C10H5 coupled with 2,4-diamino-6-(4-hydroxyanilino)-1,3,5-triazine and treated with aq. HCHO gave a dye which dyed cotton a wetfast reddish yellow shade. ANSWER 2 OF 2 CAPLUS COPYRIGHT 2003 ACS on STN L9 1962:54459 CAPLUS ANDN56:54459 OREF 56:10419h-i,10420a-g Coating of articles, especially fibers and fabrics, TIwith fibrous boehmite Bugosh, John IN E. I. du Pont de Nemours & Co. PAPatent DT Unavailable LΑ APPLICATION NO. DATE PATENT NO. KIND DATE _____ 19591130 US 3013901 US 19611219 PΙ A process is described for coating neg. charged solids with fibrous AB boehmite (I) and for use of I as an anchoring agent for subsequently applied topcoat materials. I was prepd. by adding 539.4 g. Et20-extd. Al dust to 4000 g. H2O and 965.76 g. AlCl3.6H2O with agitation for 10 hrs. at 75-95.degree. to form a slightly opalescent soln., pH 3.5, Al203:Cl ratio 1:1, which was dild. to 2% I and autoclaved at 160.degree. to form a stable, opalescent sol, pH 3.78. The sol, was deionized to pH 5.51 with an anion exchanger, giving an 11:1 Al203:Cl ratio with I having a sp. surface area of 400 sq. m./g. Regenerated cellulosic materials, solid org. plastics, paper, fibers, woven fabrics, and polyurethan foams are suitable base materials. Thus, unsized paper and wool gabardine fabric were each treated with a 2% I sol, and a 1% aq. dispersion of perfluorooctanoic acid. They had excellent H2O, oil, and grease repellency. Cotton fabric treated with 1 and Na stearate had excellent

H2O repellency after dry cleaning and soap-soln. washing. I treated Dacron, Orlon, fibrous glass, cotton cloth, and unsized filter paper

treated with acidic, mordant, or direct dyes retained the dyes after boiling; untreated fabrics lost their color. Sheepskins treated with I and NH4 perfluorooctanoate gave soft, pliable skins which were H2O- and grease-repellent, dry cleanable, and dyeable. Hydrophobic effects were obtained by spray coating bricks, concrete, plaster, wood, and terrazzo with I and then washing with Na stearate, polycarboxylic acid resins, or polymerizable carboxylic and sulfonic acid copolymers. Treatment of yarn and staple of fibers of polymers and copolymers of acrylonitrile, linear polyamides, and linear polyesters with I gave antistatic protection, increased ease of processing, and improved dyeing. Orlon jersey, treated with I, then scoured with a mixt. of 1% of the Na salt of the sulfate of the condensation product of ethylene oxide and oleyl alc. plus 1% Na3HP2O7 was more stretch resistant and less pilled than untreated Orlon. 1-treated Aerilan was more stretch-resistant and had better antistatic properties than the control. Treated Orlon upholstery fabric had better wear resistance than untreated fabric. I-treated carpet did not burn after ignition with hexamethylenetetramine, while untreated carpet was completely consumed. 1-treated tufted, undyed, and dyed carpet had better soil ratings then untreated carpet. Immersion of Al sheet which had been cleaned with NaOH and HNO3 in 1, then in a colloidal silica sol, and baking at 150-350.degree. gave a hard, adherent, abrasion-resistant coating. I was used to fix acid, mordant, or direct dyes to Al, for decorative effects, and to neg. charge colloidal graphite for improved lubricity and decreased tendency to sieze under high pressure. Al and stainless-steel panels treated with I were permanently dyed with red dye. I also imparted improved resistance to breakage to glass. Reflectance measurements on cardboard, pine boards, and plywood coated with alkyd-resin white paint and I, after application of C black, indicated excellent soil proofing. A film made by pressing I milled with granular polyethylene at 8000 lb./sq. in. and 145.degree. between At foil adhered strongly, while polyethylene film made without I did not. Addn. of 0.5-6% I powder to tobacco gave improved taste and mildness to cigarets with less tar in the smoke: cleaner and drier pipe bowls resulted from the use of I in pipe tobacco. Natural rubber-butadiene/styrene blend foams treated with 0.6-3.1% I had 40-121% higher load-bearing capacities than untreated foams. Polyurethan foams impregnated with a 2.8% I dispersion, then with polysilicic acid, poly(vinyl alc.), etc., had wicking rates of 0.125-1.0 in./sec. vs. 0.0625 in./sec. for untreated foams. These foams retained their hydrophilic character after extensive washing with detergents; they were useful as sponges.